

AMENDMENTS TO THE CLAIMS:

The following is a complete listing of the claims and reflects all changes currently being made to the claims. This listing supersedes all earlier versions and all earlier listings of the claims.

1. (Currently Amended) A solid pharmaceutical dosage form comprising caffeine, a disintegrant selected from the group consisting of sodium starch glycolate, ~~crosslinked polyvinylpyrrolidone~~, crosslinked carboxymethylcellulose, and mixtures thereof, and a cephalagic, wherein said caffeine is in the form of uncoated particles having an average particle size of about 70 to 600 microns.

2. (Previously Presented) The dosage form of claim 1, wherein the cephalagic is selected from analgesics, non-steroidal anti-inflammatory drugs, decongestants, and antihistamines.

3. (Previously Presented) The dosage form of claim 1, wherein the cephalagic is selected from the group consisting of acetaminophen, ibuprofen, ketoprofen, chlorpheniramine, diphenhydramine, and doxycyclamine.

4. (Previously Presented) The dosage form of claim 1, wherein the cephalagic is in the form of a granulation.

5. (Previously Presented) The dosage form of claim 4, wherein the granulation has an average particle size of about 100 to 400 microns.

6. (Previously Presented) The dosage form of claim 1 in the form of a directly compressed tablet.

7. (Previously Presented) The dosage form of claim 1, wherein at least 95 % of the caffeine dissolves within 5 minutes, when measured by USP, Type II Apparatus (Paddles) set at 50 rpm.

8. (Withdrawn) A process for making a solid, pharmaceutical dosage form, which comprises dry blending caffeine and a cephalagic into a blend, wherein said caffeine is in the form of uncoated particles having an average particle size of about 70 to 600 microns, and compressing the blend.

9. (Withdrawn) The process of claim 8, wherein the cephalagic is selected from the group consisting of acetaminophen, ibuprofen, ketoprofen, chlorpheniramine, diphenhydramine, and doxylamine.

10. (Withdrawn) The process of claim 8, wherein the cephalagic is in the form of a granulation.

11. (Withdrawn) The process of claim 10, wherein the granulation has an average particle size of about 100 to 400 microns.

12. (Withdrawn) The process of claim 8, wherein at least 95 % of the caffeine in the compressed blend dissolves within 5 minutes, when measured by USP, Type II Apparatus (Paddles) set at 50 rpm.

13. (Withdrawn) A compressed tablet made by the process of claim 8.